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NEWS 2 DEC 01 ChemPort single article sales feature unavailable
NEWS 3 JUN 01 CAS REGISTRY Source of Registration (SR) searching
enhanced on STN
NEWS 4 JUN 26 NUTRACEUT and PHARMAML no longer updated
NEWS 5 JUN 29 IMSCOPROFILE now reloaded monthly
NEWS 6 JUN 29 EFFULL adds Simultaneous Left and Right Truncation
(SLART) to AB, MCLM, and TI fields
NEWS 7 JUL 09 PATDPAFULL adds Simultaneous Left and Right
Truncation (SLART) to AB, CLM, MCLM, and TI fields
NEWS 8 JUL 14 USGENE enhances coverage of patent sequence location
(PSL) data
NEWS 9 JUL 27 CA/CAPLUS enhanced with new citing references
NEWS 10 JUL 16 GBFULL adds patent backfile data to 1855
NEWS 11 JUL 21 USGENE adds bibliographic and sequence information
NEWS 12 JUL 28 EFFULL adds first-page images and applicant-cited
references
NEWS 13 JUL 28 INPADOCDB and INPAFAMDB add Russian legal status data
NEWS 14 AUG 08 Improve STN by completing a survey and be entered to
win a gift card
NEWS 15 AUG 10 Time limit for inactive STN sessions doubles to 40
minutes
NEWS 16 AUG 17 CAS REGISTRY, the Global Standard for Chemical
Research, Approaches 50 Millionth Registration
Milestone
NEWS 17 AUG 18 COMPENDEX indexing changed for the Corporate Source
(CS) field

NEWS EXPRESS MAY 26 09 CURRENT WINDOWS VERSION IS V8.4,
AND CURRENT DISCOVER FILE IS DATED 06 APRIL 2009.

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 * See NEWS 14 for details or go directly to the survey at: *
 * <http://www.zoomerang.com/Survey/?p=WEB229H4S8Q5UL> *
 *

 * * * * * STN Columbus * * * * *

FILE 'HOME' ENTERED AT 12:52:36 ON 18 AUG 2009

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=> file reg
COST IN U.S. DOLLARS                SINCE FILE      TOTAL
                                     ENTRY      SESSION
FULL ESTIMATED COST                0.44          0.44
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FILE 'REGISTRY' ENTERED AT 12:53:40 ON 18 AUG 2009
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STRUCTURE FILE UPDATES: 17 AUG 2009 HIGHEST RN 1174495-28-3
 DICTIONARY FILE UPDATES: 17 AUG 2009 HIGHEST RN 1174495-28-3

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TSCA INFORMATION NOW CURRENT THROUGH June 26, 2009.

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 experimental property data in the original document. For information
 on property searching in REGISTRY, refer to:

<http://www.cas.org/support/stngen/stndoc/properties.html>

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 Uploading C:\Program Files\STNEXP\Queries\10591722b.str



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chain nodes :
1  2  3  4  5  10 19 20
ring nodes :
11 14 15 16 17 18
chain bonds :
1-3  2-5  3-4  4-5  4-10 10-11 16-19 19-20
ring bonds :
11-14 11-18 14-15 15-16 16-17 17-18
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exact/norm bonds :
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exact bonds :
10-11
normalized bonds :
11-14 11-18 14-15 15-16 16-17 17-18

Match level :
1:Atom 2:Atom 3:CLASS 4:CLASS 5:CLASS 10:CLASS 11:Atom 14:Atom 15:Atom
16:Atom 17:Atom 18:Atom 19:CLASS 20:CLASS

L1 STRUCTURE UPLOADED

=> s l1 sss full
FULL SEARCH INITIATED 12:56:15 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 3619493 TO ITERATE

43.0% PROCESSED	1556648 ITERATIONS	394 ANSWERS
50.6% PROCESSED	1829879 ITERATIONS	395 ANSWERS
55.0% PROCESSED	1992259 ITERATIONS	396 ANSWERS
55.3% PROCESSED	2000000 ITERATIONS	396 ANSWERS
INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)		
SEARCH TIME: 00.00.53		

FULL FILE PROJECTIONS: ONLINE **INCOMPLETE**
BATCH **INCOMPLETE**
PROJECTED ITERATIONS: 3619493 TO 3619493
PROJECTED ANSWERS: 636 TO 796

L2 396 SEA SSS FUL L1

=> file capl		
COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	188.28	188.72

FILE 'CAPLUS' ENTERED AT 12:57:31 ON 18 AUG 2009
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FILE COVERS 1907 - 18 Aug 2009 VOL 151 ISS 8
FILE LAST UPDATED: 17 Aug 2009 (20090817/ED)
REVISED CLASS FIELDS (/NCL) LAST RELOADED: Jun 2009

USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Jun 2009

CAPLUS now includes complete International Patent Classification (IPC) reclassification data for the second quarter of 2009.

CAS Information Use Policies apply and are available at:

<http://www.cas.org/legal/infopolicy.html>

This file contains CAS Registry Numbers for easy and accurate substance identification.

The ALL, BIB, MAX, and STD display formats in the CA/CAPLUS family of databases have been updated to include new citing references information. This enhancement may impact record import into database management software. For additional information, refer to NEWS 9.

=> s l2

L3 16 L2

=> d l3 1-16 ibib hitstr

L3 ANSWER 1 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2009:875995 CAPLUS

DOCUMENT NUMBER: 151:115083

TITLE: Method using lifespan-altering compounds for altering the lifespan of eukaryotic organisms, and screening for such compounds

INVENTOR(S): Goldfarb, David Scott

PATENT ASSIGNEE(S): University of Rochester, USA

SOURCE: U.S. Pat. Appl. Publ., 57pp.

CODEN: USXXCO

DOCUMENT TYPE: Patent

LANGUAGE: English

PATENT INFORMATION:

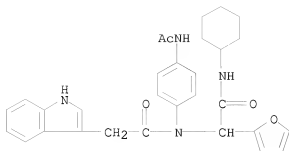
PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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US 20090163545 A1		20090625	US 2008-XQ341615	20081222
PRIORITY APPLN. INFO.:			US 2007-16362P	20071221
			US 2008-23801P	20080125

IT 1032762-38-1

RL: PAC (Pharmacological activity); BIOL (Biological study)
(method using lifespan-altering compds. for altering lifespan of eukaryotic organisms, and screening for such compds.)

RN 1032762-38-1 CAPLUS

CN 1H-Indole-3-acetamide, N-[4-(acetylamino)phenyl]-N-[2-(cyclohexylamino)-1-(2-furanyl)-2-oxoethyl]- (CA INDEX NAME)



L3 ANSWER 2 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2009:357199 CAPLUS

DOCUMENT NUMBER: 150:364629

TITLE: Lanthanide pyridine iminodicarboxylate chelate complexes as fluorescent markers for peptides and oligonucleotides

PATENT ASSIGNEE(S): Wallac Oy, Finland

SOURCE: Ger. Gebrauchsmusterschrift, 7pp.

CODEN: GGXXFR

DOCUMENT TYPE: Patent

LANGUAGE: German

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 202008013315	U1	20090326	DE 2008-202008013315	20081007
PRIORITY APPLN. INFO.:			FI 2007-493U	U 20071217

OTHER SOURCE(S): MARPAT 150:364629

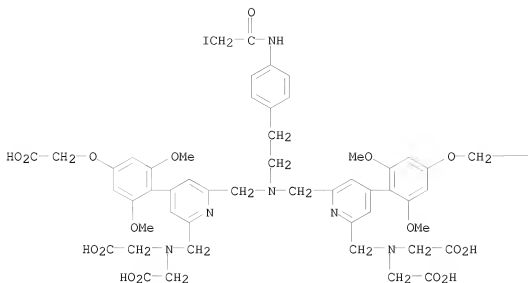
IT 1133438-12-6P 1133438-13-7P

RL: PEP (Physical, engineering or chemical process); PRPH (Prophetic); RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); PROC (Process); RACT (Reactant or reagent)

(preparation of lanthanide pyridine iminodicarboxylate chelate complexes as fluorescent markers for peptides and oligonucleotides)

RN 1133438-12-6 CAPLUS

CN INDEX NAME NOT YET ASSIGNED

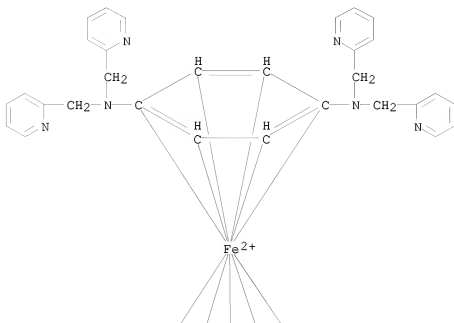


— CO_2H

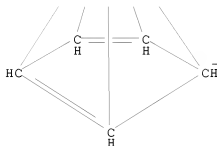
L3 ANSWER 3 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
 ACCESSION NUMBER: 2009:247359 CAPLUS
 DOCUMENT NUMBER: 150:422650
 TITLE: Ultrasound-promoted aromatic nucleophilic substitution of dichlorobenzene iron(II) complexes
 AUTHOR(S): Raouafi, Noureddine; Belhadj, Nadra; Boujlel, Khaled; Ourari, Ali; Amatore, Christian; Maisonnaute, Emmanuel; Schoellhorn, Bernd
 CORPORATE SOURCE: Departement de Chimie, Faculte des Sciences de Tunis, Universite de Tunis El Manar, Tunis, 2092, Tunisia
 SOURCE: Tetrahedron Letters (2009), 50(15), 1720-1722
 CODEN: TELEAY; ISSN: 0040-4039
 PUBLISHER: Elsevier Ltd.
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 IT 1142881-34-2P
 RL: SPN (Synthetic preparation); PREP (Preparation)
 (preparation of anilines by ultrasound-promoted aromatic nucleophilic substitution of chlorobenzene iron complexes)

RN 1142881-34-2 CAPLUS
 CN Iron(1+), (η5-2,4-cyclopentadien-1-yl) [(1,2,3,4,5,6-η)-N1,N1,N4,N4-tetrakis(2-pyridinylmethyl)-1,4-benzenediamine]- (CA INDEX NAME)

PAGE 1-A



PAGE 2-A



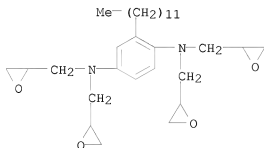
REFERENCE COUNT: 19 THERE ARE 19 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 4 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
 ACCESSION NUMBER: 2008:820982 CAPLUS
 DOCUMENT NUMBER: 149:211889
 TITLE: Vertical-alignment liquid crystal aligning agents and vertical-alignment mode liquid crystal display elements
 INVENTOR(S): Kumagaya, Tsutomu; Nishikawa, Michinori
 PATENT ASSIGNEE(S): Jsr Corporation, Japan
 SOURCE: Faming Zhuanli Shenqing Gongkai Shuomingshu, 34pp.
 CODEN: CNXXEV
 DOCUMENT TYPE: Patent
 LANGUAGE: Chinese

FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
CN 101210184	A	20080702	CN 2007-10305939	20071228
JP 2008181102	A	20080807	JP 2007-323414	20071214
KR 2008063148	A	20080703	KR 2007-138689	20071227
PRIORITY APPLN. INFO:			JP 2006-354460	A 20061228

IT 1041184-76-2
RL: TEM (Technical or engineered material use); USES (Uses)
(liquid crystal aligning agents for vertical-alignment mode liquid crystal displays)
RN 1041184-76-2 CAPLUS
CN 1,4-Benzenediamine, 2-dodecyl-N1,N1,N4,N4-tetrakis(2-oxiranylmethyl)- (CA INDEX NAME)



L3 ANSWER 5 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2008:736395 CAPLUS
DOCUMENT NUMBER: 149:79490
TITLE: Carboxamides as ion channel modulators and their preparation, pharmaceutical compositions and use in the treatment of diseases
INVENTOR(S): Galullo, Vincent; Zelle, Robert; Mazdiyasni, Hormoz; Baker, Christopher Todd; Will, Paul; Guo, Jinsong; Fensome, Andrew; Soenen, Danielle; Kern, Jeffrey Curtis; Moore, William Jay; Melenski, Edward George; Kaplan, Justin; Sabatucci, Joseph Peter
PATENT ASSIGNEE(S): Wyeth, John, and Brother Ltd., USA
SOURCE: PCT Int. Appl., 312pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2008073461	A2	20080619	WO 2007-US25416	20071211
WO 2008073461	A3	20080912		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE,

IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF,
 BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW,
 GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ,
 BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA

PRIORITY APPLN. INFO.: US 2006-874133P P 20061211
 US 2006-874152P P 20061211
 US 2006-874179P P 20061211

OTHER SOURCE(S): MARPAT 149:79490

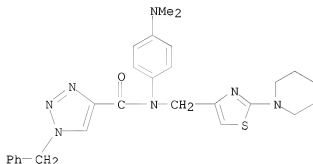
IT 1033831-47-8P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU
 (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
 (Uses)

(drug candidate; preparation of carboxamide compds. as ion channel
 modulators useful in treatment of diseases)

RN 1033831-47-8 CAPLUS

CN 1H-1,2,3-Triazole-4-carboxamide, N-[4-(dimethylamino)phenyl]-1-
 (phenylmethyl)-N-[[2-(1-piperidinyl)-4-thiazolyl]methyl]- (CA INDEX NAME)



OS.CITING REF COUNT: 2 THERE ARE 2 CAPLUS RECORDS THAT CITE THIS RECORD
 (2 CITINGS)

L3 ANSWER 6 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2008:733525 CAPLUS

DOCUMENT NUMBER: 149:53863

TITLE: Preparation of of N,N-substituted 3-aminopyrrolidine
 compounds useful as monoamines reuptake inhibitors

INVENTOR(S): Kurimura, Muneaki; Taira, Shinichi; Tomoyasu,
 Takahiro; Ito, Nobuaki; Tai, Kuninori; Takemura,
 Noriaki; Matsuzaki, Takayuki; Menjo, Yasuhiro;
 Miyamura, Shin; Sakurai, Yoji; Watabe, Akihito;
 Sakata, Yasuyo; Masumoto, Takumi; Akazawa, Kohei;
 Sugino, Haruhiko; Amada, Naoki; Ohashi, Satoshi;
 Shinohara, Tomokazu; Sasaki, Hirofumi; Morita,
 Chisako; Yamashita, Junko; Nakajima, Satoko

PATENT ASSIGNEE(S): Ohtsuka Pharmaceutical Co., Ltd., Japan
 SOURCE: Jpn. Kokai Tokkyo Koho, 221pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent
 LANGUAGE: Japanese

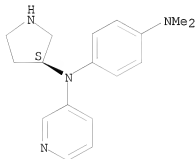
FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2008137997	A	20080619	JP 2007-292386	20071109
PRIORITY APPLN. INFO.:			JP 2006-305573	A 20061110
OTHER SOURCE(S):	MARPAT 149:53863			

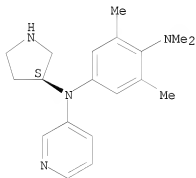
IT 914997-33-4P 914997-67-4P 914997-70-9P
 915000-91-8P 915001-14-8P
 RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU
 (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
 (Uses)
 (preparation of aminopyrrolidine compds. as monoamines reuptake inhibitors
 with sufficient therapeutic effects after short-term administration)
 RN 914997-33-4 CAPLUS
 CN 1,4-Benzenediamine, N1,N1-dimethyl-N4-3-pyridinyl-N4-(3S)-3-pyrrolidinyl-
 (CA INDEX NAME)

Absolute stereochemistry.



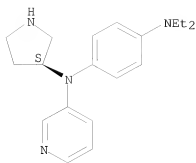
RN 914997-67-4 CAPLUS
 CN 1,4-Benzenediamine, N1,N1,2,6-tetramethyl-N4-3-pyridinyl-N4-(3S)-3-
 pyrrolidinyl- (CA INDEX NAME)

Absolute stereochemistry.



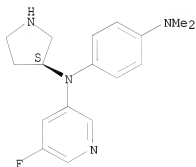
RN 914997-70-9 CAPLUS
 CN 1,4-Benzenediamine, N1,N1-diethyl-N4-3-pyridinyl-N4-(3S)-3-pyrrolidinyl-
 (CA INDEX NAME)

Absolute stereochemistry.



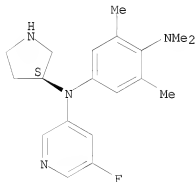
RN 915000-91-8 CAPLUS
 CN 1,4-Benzenediamine, N1-(5-fluoro-3-pyridinyl)-N4,N4-dimethyl-N1-(3S)-3-pyrrolidinyl- (CA INDEX NAME)

Absolute stereochemistry.



RN 915001-14-8 CAPLUS
 CN 1,4-Benzenediamine, N4-(5-fluoro-3-pyridinyl)-N1,N1,2,6-tetramethyl-N4-(3S)-3-pyrrolidinyl- (CA INDEX NAME)

Absolute stereochemistry.

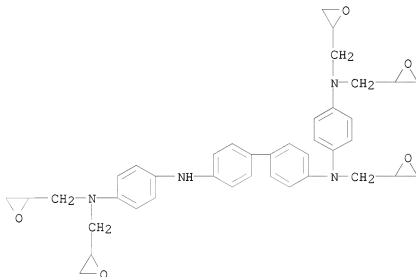


L3 ANSWER 7 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
 ACCESSION NUMBER: 2008:447206 CAPLUS
 DOCUMENT NUMBER: 148:506743
 TITLE: Liquid crystal alignment agent and liquid crystal display element
 INVENTOR(S): Yasuda, Hiroyuki; Hayashi, Eiji; Nishikawa, Michinori

PATENT ASSIGNEE(S): Jsr Corporation, Japan
 SOURCE: Faming Zhuanti Shengqing Gongkai Shuomingshu, 43pp.
 CODEN: CNXXEV
 DOCUMENT TYPE: Patent
 LANGUAGE: Chinese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
CN 101153995	A	20080402	CN 2007-10151310	20070924
KR 2008028320	A	20080331	KR 2007-96550	20070921
JP 2008107811	A	20080508	JP 2007-247604	20070925
PRIORITY APPLN. INFO.: IT 1020839-25-1P			JP 2006-261196	A 20060926

RL: SPN (Synthetic preparation); TEM (Technical or engineered material use); PREP (Preparation); USES (Uses)
 (liquid crystal alignment agent and liquid crystal display element)
 RN 1020839-25-1 CAPLUS
 CN [1,1'-Biphenyl]-4,4'-diamine, N4,N4'-bis[4-{bis(2-oxiranylmethyl)amino}phenyl]-N4-(2-oxiranylmethyl)- (CA INDEX NAME)



L3 ANSWER 8 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
 ACCESSION NUMBER: 2008:124359 CAPLUS
 DOCUMENT NUMBER: 148:191838
 TITLE: Preparation of substituted aniline derivatives as antifungal agents
 INVENTOR(S): Carr, Andrew David; Neuss, Judi Charlotte; Orchard, Michael Glen; Porter, David William
 PATENT ASSIGNEE(S): Ucb Pharma S.A., Belg.
 SOURCE: PCI Int. Appl., 11pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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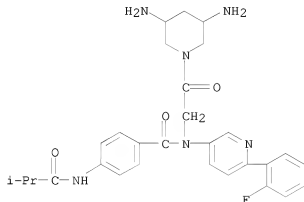
WO 2008012524 A1 20080131 WO 2007-GB2815 20070724
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RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
AU 2007279092 A1 20080131 AU 2007-279092 20070724
CA 2658913 A1 20080131 CA 2007-2658913 20070724
US 20080045497 A1 20080221 US 2007-782337 20070724
EP 2046739 A1 20090415 EP 2007-766352 20070724
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, MK, RS

PRIORITY APPLN. INFO.:

GB 2006-14677 A 20060724
GB 2006-14678 A 20060724
GB 2007-4645 A 20070309
GB 2007-4648 A 20070309
WO 2007-GB2815 W 20070724

OTHER SOURCE(S): MARPAT 148:191838

IT 1013324-35-0P
RL: PAC (Pharmacological activity); PRPH (Prophetic); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(preparation of substituted aniline derivs. as antifungal agents)
RN 1013324-35-0 CAPLUS
CN Benzamide, N-[2-(3,5-diamino-1-piperidinyl)-2-oxoethyl]-N-[6-(2-fluorophenyl)-3-pyridinyl]-4-[(2-methyl-1-oxopropyl)amino]- (CA INDEX NAME)



REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 9 OF 16 CAPLUS COPYRIGHT 2009 ACS ON STN
ACCESSION NUMBER: 2007:1364437 CAPLUS
DOCUMENT NUMBER: 148:33637
TITLE: Substituted quinolones as ATP-utilizing enzyme inhibitors and their preparation, compositions, and uses thereof

INVENTOR(S): Dickson, John K.; Chen, Ke; Hodge, Carl Nicholas
 PATENT ASSIGNEE(S): Amphora Discovery Corporation, USA
 SOURCE: PCT Int. Appl., 143pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007136592	A2	20071129	WO 2007-US11484	20070510
WO 2007136592	A3	20080228		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA CA 2652634 A1 20071129 CA 2007-2652634 20070510 US 20070287706 A1 20071213 US 2007-803140 20070510 EP 2040711 A2 20090401 EP 2007-794818 20070510 R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, MK, RS PRIORITY APPLN. INFO.: US 2006-801881P P 20060518 WO 2007-US11484 W 20070510				

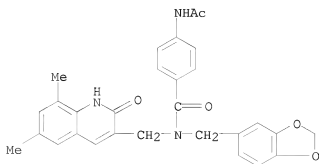
OTHER SOURCE(S): MARPAT 148:33637
 IT 958454-68-7P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; preparation of substituted quinolones as ATP-utilizing enzyme inhibitors useful in the treatment of diseases)

RN 958454-68-7 CAPLUS

CN Benzamide, 4-(acetylamino)-N-(1,3-benzodioxol-5-ylmethyl)-N-[(1,2-dihydro-6,8-dimethyl-2-oxo-3-quinolinyl)methyl]- (CA INDEX NAME)



OS.CITING REF COUNT: 1 THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD (1 CITINGS)

L3 ANSWER 10 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2007:1029651 CAPLUS

DOCUMENT NUMBER: 147:365486

TITLE: Preparation of 2-(phenylamino)thiazole derivatives as inhibitors of viral replication for the treatment of hepatitis C infection

INVENTOR(S): Zhang, Suoming; Phadke, Avinash; Wang, Xiangzhu; Liu, Cuixian

PATENT ASSIGNEE(S): Achillion Pharmaceuticals, Inc., USA

SOURCE: PCT Int. Appl., 134pp.

CODEN: PIXXD2

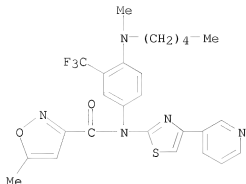
DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007103550	A2	20070913	WO 2007-US6023	20070308
WO 2007103550	A3	20071108		
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW			
RW:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA			
AU 2007223797	A1	20070913	AU 2007-223797	20070308
CA 2645072	A1	20070913	CA 2007-2645072	20070308
US 20070213301	A1	20070913	US 2007-683749	20070308
EP 1996565	A2	20081203	EP 2007-752705	20070308
R:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, MK, RS			
JP 2009529059	T	20090813	JP 2008-558416	20070308
PRIORITY APPLN. INFO.:			US 2006-780609P	P 20060308
			WO 2007-US6023	W 20070308
OTHER SOURCE(S):	CASREACT 147:365486; MARPAT 147:365486			
IT 949117-20-8P				
RL:	PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)			
	(drug candidate; preparation of (phenylamino)thiazoles as inhibitors of viral replication for treatment of hepatitis C infection)			
RN 949117-20-8	CAPLUS			
CN 3-Isoxazolecarboxamide, 5-methyl-N-[4-(methylpentylamino)-3-(trifluoromethyl)phenyl]-N-[4-(3-pyridinyl)-2-thiazolyl]-				(CA INDEX NAME)



OS.CITING REF COUNT: 1 THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD
(1 CITINGS)

L3 ANSWER 11 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2007:1028755 CAPLUS

DOCUMENT NUMBER: 147:365493

TITLE: Heterobicyclic pyrazole compounds as Met tyrosine
kinase inhibitors and their preparation and use

INVENTOR(S): Blake, James F.; Boyd, Steven Armen; Cohen, Frederick;
De Meese, Jason; Fong, Kin Chiu; Gaudino, John J.;
Kaplan, Tomas; Marlow, Allison L.; Seo, Jeongbeob;
Thomas, Allen A.; Tian, Hongqi; Young, Wendy B.

PATENT ASSIGNEE(S): Array Biopharma Inc., USA; Genentech, Inc.

SOURCE: PCT Int. Appl., 2/73 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007103308	A2	20070913	WO 2007-US5583	20070306
WO 2007103308	A3	20080207		
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW			
RW:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA			
AU 2007224020	A1	20070913	AU 2007-224020	20070306
CA 2645137	A1	20070913	CA 2007-2645137	20070306
US 20070238726	A1	20071011	US 2007-714342	20070306
EP 2001880	A2	20081217	EP 2007-752297	20070306
R:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, MT, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, MK, RS			
JP 2009529047	T	20090813	JP 2008-558335	20070306
MX 2008011220	A	20080911	MX 2008-11220	20080902
IN 2008KN03882	A	20090227	IN 2008-KN3882	20080924

NO 2008004183	A	20081124	NO 2008-4183	20081006
KR 2008110783	A	20081219	KR 2008-724415	20081006
CN 101437820	A	20090520	CN 2007-80016155	20081104
PRIORITY APPLN. INFO.:			US 2006-779805P	P 20060307
			US 2006-874832P	P 20061214
			WO 2007-US5583	W 20070306

OTHER SOURCE(S): MARPAT 147:365493

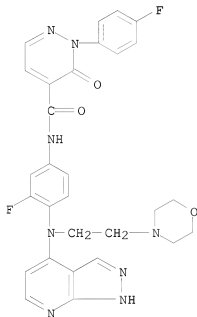
IT 949560-23-0P 949560-28-5P

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; preparation of heterobicyclic pyrazole compds. as Met tyrosine kinase inhibitors useful in the treatment of diseases)

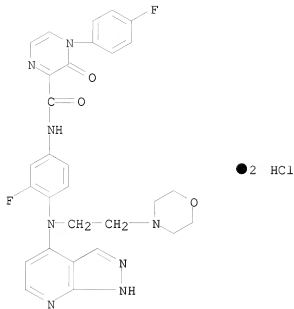
RN 949560-23-0 CAPLUS

CN 4-Pyridazinecarboxamide, N-[3-fluoro-4-[[2-(4-morpholinyl)ethyl]-1H-pyrazolo[3,4-b]pyridin-4-ylamino]phenyl]-2-(4-fluorophenyl)-2,3-dihydro-3-oxo- (CA INDEX NAME)



RN 949560-28-5 CAPLUS

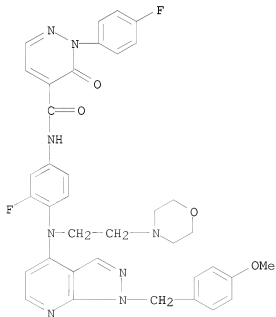
CN 2-Pyrazinecarboxamide, N-[3-fluoro-4-[[2-(4-morpholinyl)ethyl]-1H-pyrazolo[3,4-b]pyridin-4-ylamino]phenyl]-4-(4-fluorophenyl)-3,4-dihydro-3-oxo-, hydrochloride (1:2) (CA INDEX NAME)



IT 949560-27-4P, N-[4-[[1-(4-Methoxybenzyl)-1H-pyrazolo[3,4-b]pyridin-4-yl] (2-morpholinoethyl) amino]-3-fluorophenyl]-2-(4-fluorophenyl)-3-oxo-2,3-dihydropyridazine-4-carboxamide
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (intermediate; preparation of heterobicyclic pyrazole compds. as Met tyrosine kinase inhibitors useful in the treatment of diseases)

RN 949560-27-4 CAPLUS

CN 4-Pyridazinecarboxamide, N-[3-fluoro-4-[[1-[(4-methoxyphenyl)methyl]-1H-pyrazolo[3,4-b]pyridin-4-yl] [2-(4-morpholinyl)ethyl]amino]phenyl]-2-(4-fluorophenyl)-2,3-dihydro-3-oxo- (CA INDEX NAME)

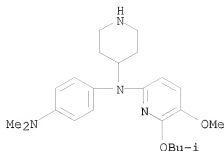


OS.CITING REF COUNT: 1 THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD
(1 CITINGS)

L3 ANSWER 12 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
ACCESSION NUMBER: 2006:1338137 CAPLUS
DOCUMENT NUMBER: 146:81773
TITLE: Preparation of N-substituted diarylamine analogs as
phosphodiesterase 4 inhibitors
INVENTOR(S): Talamas, Francisco Xavier; Caroon, Joan Marie; Dunn,
Robert; Hopper, Allen; Kuester, Eric; Schumacher,
Richard; Tehim, Ashok
PATENT ASSIGNEE(S): Memory Pharmaceuticals Corporation, USA; F.
Hoffmann-La Roche A.-G.
SOURCE: PCT Int. Appl., 114 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006135828	A2	20061221	WO 2006-US22655	20060609
WO 2006135828	A3	20070426		
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
RW:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AP, EA, EP, OA			
AU 2006257863	A1	20061221	AU 2006-257863	20060609
CA 2611562	A1	20061221	CA 2006-2611562	20060609
US 20070049611	A1	20070301	US 2006-449868	20060609
EP 1888528	A2	20080220	EP 2006-784743	20060609
R:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, AL, BA, HR, MK, YU			
JP 2008543781	T	20081204	JP 2008-515998	20060609
US 20090118270	A1	20090507	US 2008-329214	20081205
PRIORITY APPLN. INFO.:			US 2005-689060P	P 20050610
			US 2006-449868	A1 20060609
			WO 2006-US22655	W 20060609

OTHER SOURCE(S): MARPAT 146:81773
IT 917098-71-6P, N-(6-Isobutoxy-5-methoxy-pyridin-2-yl)-N',N'-dimethyl-N-(piperidin-4-yl)benzene-1,4-diamine
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(drug candidate; preparation of N-substituted diarylamine analogs as phosphodiesterase 4 inhibitors for treating cognition disorders, inflammation, and other disorders)
RN 917098-71-6 CAPLUS
CN 1,4-Benzenediamine, N1-[5-methoxy-6-(2-methylpropoxy)-2-pyridinyl]-N4,N4-dimethyl-N1-4-piperidinyl- (CA INDEX NAME)



OS.CITING REF COUNT: 1 THERE ARE 1 CAPLUS RECORDS THAT CITE THIS RECORD
(1 CITINGS)
REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS
RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 13 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2006:1206758 CAPLUS

DOCUMENT NUMBER: 145:505325

TITLE: Preparation of of N,N-substituted 3-aminopyrrolidine

compounds useful as monoamines reuptake inhibitors
INVENTOR(S): Kurimura, Muneaki; Taira, Shinichi; Tomoyasu,
Takahiro; Ito, Nobuaki; Tai, Kuninori; Takemura,
Noriaki; Matsuzaki, Takayuki; Menjo, Yasuhiro;
Miyamura, Shin; Sakurai, Yohji; Watanabe, Akihito;
Sakata, Yasuyo; Masumoto, Takumi; Akazawa, Kohei;
Sugino, Haruhiko; Amada, Naoki; Ohashi, Satoshi;
Shinohara, Tomochi; Sasaki, Hirofumi; Morita,
Chisako; Yamashita, Junko; Nakajima, Satoko

PATENT ASSIGNEE(S): Otsuka Pharmaceutical Co., Ltd., Japan

SOURCE: PCT Int. Appl., 260pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006121218	A1	20061116	WO 2006-JP309988	20060512
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
AU 2006244851	A1	20061116	AU 2006-244851	20060512
CA 2608184	A1	20061116	CA 2006-2608184	20060512
EP 1881975	A1	20080130	EP 2006-756356	20060512
R: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR				
JP 2008540329	T	20081120	JP 2007-552427	20060512
IN 2007DN08276	A	20071123	IN 2007-DN8276	20071026
CN 101175748	A	20080507	CN 2006-80016402	20071112

MX 2007014252	A	20080122	MX 2007-14252	20071113
KR 2008008423	A	20080123	KR 2007-729022	20071212
US 20090088406	A1	20090402	US 2008-914183	20080929
PRIORITY APPLN. INFO.:			JP 2005-141230	A 20050513
			WO 2006-JP309988	W 20060512

OTHER SOURCE(S): MARPAT 145:505325

IT 914997-33-4P 914997-67-4P 914997-70-9P
 915000-91-8P 915001-14-8P

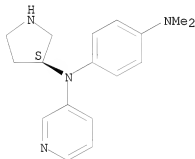
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU
 (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES
 (Uses)

(preparation of aminopyrrolidine compds. as monoamines reuptake inhibitors
 with sufficient therapeutic effects after short-term administration)

RN 914997-33-4 CAPLUS

CN 1,4-Benzenediamine, N1,N1-dimethyl-N4-3-pyridinyl-N4-(3S)-3-pyrrolidinyl-
 (CA INDEX NAME)

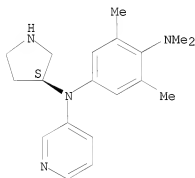
Absolute stereochemistry.



RN 914997-67-4 CAPLUS

CN 1,4-Benzenediamine, N1,N1,2,6-tetramethyl-N4-3-pyridinyl-N4-(3S)-3-
 pyrrolidinyl- (CA INDEX NAME)

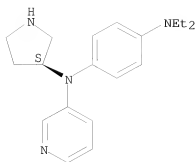
Absolute stereochemistry.



RN 914997-70-9 CAPLUS

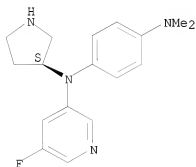
CN 1,4-Benzenediamine, N1,N1-diethyl-N4-3-pyridinyl-N4-(3S)-3-pyrrolidinyl-
 (CA INDEX NAME)

Absolute stereochemistry.



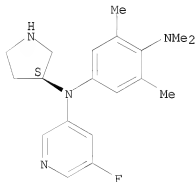
RN 915000-91-8 CAPLUS
 CN 1,4-Benzenediamine, N1-(5-fluoro-3-pyridinyl)-N4,N4-dimethyl-N1-(3S)-3-pyrrolidinyl- (CA INDEX NAME)

Absolute stereochemistry.



RN 915001-14-8 CAPLUS
 CN 1,4-Benzenediamine, N4-(5-fluoro-3-pyridinyl)-N1,N1,2,6-tetramethyl-N4-(3S)-3-pyrrolidinyl- (CA INDEX NAME)

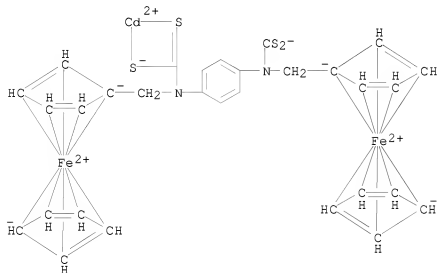
Absolute stereochemistry.



REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 14 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
 ACCESSION NUMBER: 2006:109700 CAPLUS
 DOCUMENT NUMBER: 145:431181
 TITLE: Synthesis and thermal decomposition of cadmium

AUTHOR(S): dithiocarbamate complexes
 CORPORATE SOURCE: Thammakan, Nirawan; Somsook, Ekasith
 Department of Chemistry, Faculty of Science, Mahidol
 University, Bangkok, 10400, Thailand
 SOURCE: Materials Letters (2006), 60(9-10), 1161-1165
 CODEN: MLETDJ; ISSN: 0167-577X
 PUBLISHER: Elsevier B.V.
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 OTHER SOURCE(S): CASREACT 145:431181
 IT 911824-19-6P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT
 (Reactant or reagent)
 (preparation and thermal decomposition of cadmium(II) benzyl- and
 ferrocenylmethyl-substituted benzenedithiocarbamate polymeric
 complexes)
 RN 911824-19-6 CAPLUS
 CN Cadmium, [[[(dithiocarboxy-κS,κS') [4-
 [(dithiocarboxy)(ferrocenylmethyl)amino]phenyl]amino]methyl]ferrocenato(2-
)]- (9CI) (CA INDEX NAME)



OS.CITING REF COUNT: 2 THERE ARE 2 CAPLUS RECORDS THAT CITE THIS RECORD
 (2 CITINGS)
 REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS
 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L3 ANSWER 15 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN
 ACCESSION NUMBER: 2005:1075811 CAPLUS
 DOCUMENT NUMBER: 143:367523
 TITLE: Preparation of monosaccharide derivatives as
 anti-inflammatory agents
 INVENTOR(S): Sattigeri, Viswajanani Jitendra; Arora, Sudershan K.;
 Salman, Mohammad; Palle, Venkata P.; Yadav, Gyan
 Chand; Tanwar, Madan Pal; Mukherjee, Ashis; Narayanan,
 Ramamurthy; Rauf, Abdul Rehman Abdul; Naik, Keshav
 Prabhakar; Soni, Ajay; Ray, Abhijit; Shirumalla, Raj
 Kumar; Mookhtiar, Kasim Abbas
 PATENT ASSIGNEE(S): Ranbaxy Laboratories Limited, India
 SOURCE: PCT Int. Appl., 185 pp.
 CODEN: PIXXD2

DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005092907	A2	20051006	WO 2005-IB803	20050329
WO 2005092907	A3	20060427		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW

RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

PRIORITY APPLN. INFO.: US 2004-556936P P 20040326

OTHER SOURCE(S): CASREACT 143:367523; MARPAT 143:367523

IT 1043943-91-4

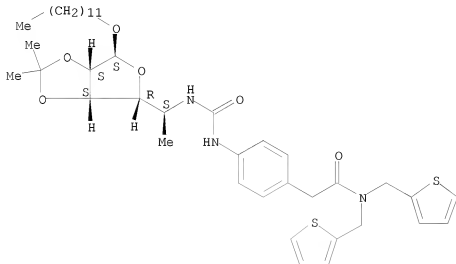
RL: PRPH (Prophetic)

(Preparation of monosaccharide derivatives as anti-inflammatory agents)

RN 1043943-91-4 CAPLUS

CN INDEX NAME NOT YET ASSIGNED

Relative stereochemistry.



L3 ANSWER 16 OF 16 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2002:487497 CAPLUS

DOCUMENT NUMBER: 137:78952

TITLE: Preparation of substituted imidazoles, pyrazoles and amides as high affinity C5a receptor modulators

INVENTOR(S): Thurkauf, Andrew; Zhang, Xiaoyan; He, Xia-Shu; Zhao, He; Peterson, John; Maynard, George; Ohliger, Robert

PATENT ASSIGNEE(S): Neurogen Corporation, USA

SOURCE: PCT Int. Appl., 609 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002049993	A2	20020627	WO 2000-US26816	20000929
WO 2002049993	A3	20030220		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
CA 2420215	A1	20020627	CA 2000-2420215	20000929
AU 2000076225	A	20020701	AU 2000-76225	20000929
EP 1322309	A2	20030702	EP 2000-965522	20000929
EP 1322309	B1	20080813		
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL				
ZA 2003001160	A	20040212	ZA 2003-1160	20000929
BR 2000017338	A	20040427	BR 2000-17338	20000929
JP 2004525873	T	20040826	JP 2002-551496	20000929
AU 2000276225	B2	20080710	AU 2000-276225	20000929
AT 404553	T	20080815	AT 2000-965522	20000929
NO 2003001370	A	20030530	NO 2003-1370	20030326
MX 2003002788	A	20041213	MX 2003-2788	20030328
PRIORITY APPLN. INFO.:				
			US 2000-227454P	P 20000823
			US 1999-156390P	P 19990928
			US 2000-202749P	P 20000508
			US 2000-212449P	P 20000616
			US 2000-221787P	P 20000731
			US 2000-224036P	P 20000809
			WO 2000-US26816	W 20000929

OTHER SOURCE(S): MARPAT 137:78952

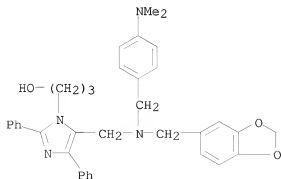
IT 1106056-27-2

RL: PRPH (Prophetic)

(Preparation of substituted imidazoles, pyrazoles and amides as high affinity C5a receptor modulators)

RN 1106056-27-2 CAPLUS

CN 1H-Imidazole-1-propanol, 5-[[[1,3-benzodioxol-5-ylmethyl][4-(dimethylamino)phenyl]methyl]amino]methyl]-2,4-diphenyl- (CA INDEX NAME)



OS.CITING REF COUNT: 9 THERE ARE 9 CAPLUS RECORDS THAT CITE THIS RECORD
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